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Life Sciences Reporting Summary

Nature Research wishes to improve the reproducibility of the work that we publish. This form is intended for publication with all accepted life science papers and provides structure for consistency and transparency in reporting. Every life science submission will use this form; some list items might not apply to an individual manuscript, but all fields must be completed for clarity.

For further information on the points included in this form, see Reporting Life Sciences Research. For further information on Nature Research policies, including our data availability policy, see Authors & Referees and the Editorial Policy Checklist.

Experimental design

1. Sample size

Describe how sample size was determined.

Sample size was calculated by survey/census source and was taken to be the sum of those sampled, after cleaning the data. The number of latitude-longitude referenced point data clusters, the number of areal polygons, and the total number of individuals from each source may be found in Supplementary Table 2.

2. Data exclusions

Describe any data exclusions.

Select data sources that were identified to contain years of education within the geographic area of interest were excluded for the following reasons: missing survey weights for areal data, missing gender variable, incomplete sampling (e.g., only a specific age range), or untrustworthy data (as determined by the survey administrator or by inspection). Within each source, administrative units with a sample size of one were excluded.

3. Replication

Describe whether the experimental findings were reliably reproduced.

4. Randomization

Describe how samples/organisms/participants were allocated into experimental groups.

5. Blinding

Describe whether the investigators were blinded to group allocation during data collection and/or analysis.

This is an observational study using many years of survey data and could be replicated.

The data in our study predominantly comes from surveys with randomized survey designs. As an observational mapping project, there were no experimental groups.

Blinding was not relevant to this study, as it was an observational study using survey data.

Note: all studies involving animals and/or human research participants must disclose whether blinding and randomization were used.

For all figures and tables that use statistical methods, conf Methods section if additional space is needed).	firm that the following items are present in relevant figure legends (or in the
n/a Confirmed	
The exact sample size (n) for each experimental group/cc	ondition, given as a discrete number and unit of measurement (animals, litters, cultures, etc.)
A description of how samples were collected, noting sample was measured repeatedly	whether measurements were taken from distinct samples or whether the same
A statement indicating how many times each experin	ment was replicated
The statistical test(s) used and whether they are one complex techniques should be described in the Meth	- or two-sided (note: only common tests should be described solely by name; more nods section)
A description of any assumptions or corrections, such	h as an adjustment for multiple comparisons
The test results (e.g. P values) given as exact values w	whenever possible and with confidence intervals noted
A clear description of statistics including central tend	lency (e.g. median, mean) and variation (e.g. standard deviation, interquartile range)
Clearly defined error bars	
See the web collection on stati.	stics for biologists for further resources and guidance.
▶ Software	
Policy information about availability of computer code	
7. Software	
Describe the software used to analyze the data in this study.	The models were all fit using R version 3.3.2. The main statistical space-time Gaussian process regression models were fit using R-INLA version 0.0-1440400394.
	central to the paper but not yet described in the published literature, software must be made ourage code deposition in a community repository (e.g. GitHub). <i>Nature Methods</i> guidance for information on this topic.
► Materials and reagents	
Policy information about availability of materials	
8. Materials availability	
Indicate whether there are restrictions on availability of unique materials or if these materials are only available for distribution by a for-profit company.	No unique materials were used.
9. Antibodies	
Describe the antibodies used and how they were validated for use in the system under study (i.e. assay and species).	No antibodies were used.
10. Eukaryotic cell lines	
a. State the source of each eukaryotic cell line used.	No eukaryotic cell lines were used.
b. Describe the method of cell line authentication used.	No eukaryotic cell lines were used.
 Report whether the cell lines were tested for mycoplasma contamination. 	No eukaryotic cell lines were used.
d. If any of the cell lines used are listed in the database of commonly misidentified cell lines maintained by ICLAC, provide a scientific rationale for their use.	No commonly misidentified cell lines were used.
▶ Animals and human research participant	ts
Policy information about studies involving animals; when repo	orting animal research, follow the ARRIVE guidelines
11. Description of research animals	
Provide details on animals and/or animal-derived materials used in the study.	No animals were used.

6. Statistical parameters

Policy information about studies involving human research participants

12. Description of human research participants

Describe the covariate-relevant population characteristics of the human research participants.

The study did not involve human research participants, as all data was obtained from secondary sources.